

GenCore version 5.1.4_p5_4578
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OM nucleic - nucleic search, using sw model

Run on: March 10, 2003, 21:31:17 ; Search time 142.984 Seconds
(without alignments)
488.248 Million cell updates/sec

Title: us-09-913-524-34

Perfect score: 31
Sequence: 1 atcattgtccctctgtgctatcgcaact 31

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_101002:*
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2: /SID22/gcgdata/geneseq/geneseq-emb1/NA1981.DAT.*
3: /SID22/gcgdata/geneseq/geneseq-emb1/NA1982.DAT.*
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18: /SID22/gcgdata/geneseq/geneseq-emb1/NA1997.DAT.*
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21: /SID22/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
22: /SID22/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SID22/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SID22/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query Match	Score	Length	DB ID	Description
C 1	31	100.0	405	24	ABL64291 Stomach cancer rel
2	31	100.0	425	7	AAN60429 Sequence encoding
3	31	100.0	1620	22	AAH57530 Human pancreas cel
4	31	100.0	1630	8	AAN70315 Sequence encoding
5	31	100.0	1700	22	AAO60008 Angiotensin conver
6	31	100.0	1873	11	AAQ01648 BUF-3 gene for hum
7	31	100.0	4068	24	ABL62912 Breast cancer rela
8	31	100.0	4068	24	ABL63117 Breast cancer rela
9	31	100.0	4068	24	ABL64417 Stomach cancer rel

10	31	100.0	14416	22	AAO5491 Human reproductive
11	31	100.0	14416	23	ABL98344 Human testicular a
12	26.2	84.5	958	7	AAN60427 Sequence encoding
13	23	74.2	3588	8	AAN70317 Sequence encoding
14	21.4	69.0	1687	12	AAQ10890 Encodes Xenopus Bo
15	20.6	66.5	9662	22	AAO7086 Human reproductive
16	20.6	66.5	17705	22	AAO7085 Human reproductive
17	19.8	63.9	406	16	AAQ89258 BMP-6 encoding DNA
18	19.8	63.9	406	19	AAV24032 Bone morphogenetic
19	19.8	63.9	497	20	AAV9382 cDNA encoding bone
20	19.8	63.9	894	13	AAQ23678 Encodes C-terminal
21	19.8	63.9	894	14	AAQ37567 BMP-6 coding seque
22	19.8	63.9	894	20	AAV9261 DNA encoding carbo
23	19.8	63.9	1350	22	AAH76483 BMP 7/6 glu nucleo
24	19.8	63.9	1353	22	AAH76484 BMP 7/6 myc nucleo
25	19.8	63.9	1362	22	AAH76481 BMP 4/6 glu nucleo
26	19.8	63.9	2385	23	AAV65056 DNA encoding novel
27	19.8	63.9	2385	23	AAV78659 DNA encoding novel
28	19.8	63.9	2385	23	AAV81638 DNA encoding novel
29	19.8	63.9	2471	23	AAV71026 DNA encoding novel
30	19.8	63.9	2923	11	AAQ06173 Human Bone Morphog
31	19.8	63.9	2923	13	AAQ32855 BMP6. Rattus ratt
32	19.8	63.9	2923	14	AAQ37568 Human BMP-6 coding
33	19.8	63.9	2923	14	AAQ41294 Human BMP-6 gene.
34	19.8	63.9	3184	23	AAV87074 DNA encoding novel
35	19.8	63.9	5651	21	AAV77516 Human ORFX ORF3071
36	19.8	63.9	5801	22	AAV22778 Human cDNA encoding
C 37	19.8	63.9	31169	22	AAV41761 Genomic sequence #
C 38	19.8	63.9	31169	22	AAV75191 Human immune/haema
C 39	19.8	63.9	31169	23	ABK44029 Genomic DNA encodi
40	19.6	63.2	167343	24	ABL64403 Stomach cancer rel
41	19.6	63.2	167343	24	ABL67239 Thyroid cancer rel
C 42	19	61.3	1548	19	AAV66850 Chlamydia 16S ribo
C 43	19	61.3	2493	23	AAV88122 DNA encoding novel
C 44	19	61.3	5741	12	AAQ14939 Bacterial amylase
45	18.8	60.6	99	16	AAQ82942 Partial coding seq

ALIGNMENTS

RESULT 1
ABL64291/c
ID ABL64291 standard; DNA: 405 BP.
XX
AC ABL64291;
XX
DT 15-MAY-2002 (first entry)
XX
DE Stomach cancer related gene sequence SEQ ID NO:2628.
XX
KW Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
KW stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
KW cytostatic; gene therapy; antineoplastic; Wilms' tumour; adenocarcinoma;
KW gene; ds.
XX
OS Homo sapiens.
XX
PN WC200194629-A2.
XX
PD 13-DEC-2001.
XX
PF 30-MAY-2001; 2001WO-US10838.
XX
PR 05-JUN-2000; 2000US-209473P.
PR 05-JUN-2000; 2000US-209531P.
PR 18-SEP-2000; 2000US-233133P.
PR 18-SEP-2000; 2000US-233617P.
PR 20-SEP-2000; 2000US-234009P.
PR 20-SEP-2000; 2000US-234034P.
PR 20-SEP-2000; 2000US-234052P.
PR 22-SEP-2000; 2000US-234509P.
PR 22-SEP-2000; 2000US-234567P.

PR 25-SEP-2000; 2000US-234923P.
 PR 25-SEP-2000; 2000US-234924P.
 PR 25-SEP-2000; 2000US-235077P.
 PR 25-SEP-2000; 2000US-235082P.
 PR 25-SEP-2000; 2000US-235134P.
 PR 25-SEP-2000; 2000US-235280P.
 PR 26-SEP-2000; 2000US-235537P.
 PR 26-SEP-2000; 2000US-235538P.
 PR 27-SEP-2000; 2000US-235711P.
 PR 27-SEP-2000; 2000US-235720P.
 PR 27-SEP-2000; 2000US-235840P.
 PR 27-SEP-2000; 2000US-235863P.
 PR 28-SEP-2000; 2000US-236028P.
 PR 28-SEP-2000; 2000US-236032P.
 PR 28-SEP-2000; 2000US-236033P.
 PR 28-SEP-2000; 2000US-236034P.
 PR 28-SEP-2000; 2000US-236109P.
 PR 28-SEP-2000; 2000US-236111P.
 PR 29-SEP-2000; 2000US-236842P.
 PR 29-SEP-2000; 2000US-236891P.
 PR 02-OCT-2000; 2000US-237172P.
 PR 02-OCT-2000; 2000US-237173P.
 PR 02-OCT-2000; 2000US-237278P.
 PR 02-OCT-2000; 2000US-237294P.
 PR 02-OCT-2000; 2000US-237295P.
 PR 02-OCT-2000; 2000US-237316P.
 PR 03-OCT-2000; 2000US-237425P.
 PR 03-OCT-2000; 2000US-237598P.
 PR 03-OCT-2000; 2000US-237604P.
 PR 03-OCT-2000; 2000US-237606P.
 PR 03-OCT-2000; 2000US-237608P.
 PR 01-NOV-2000; 2000US-244867P.
 PR 01-NOV-2000; 2000US-245084P.
 XX
 PA (AVAL-) AVALON PHARM.

XX Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;

PI Soppet DR, Weaver Z;

XX WPI: 2002-188264/24.

XX Screening for anti-neoplastic agent involves exposing cells to a

PI chemical agent to be tested for anti-neoplastic activity, and

PT determining a change in expression of a gene of a signature gene set

XX Claim 1; SEQ ID 2628; 44pp; English.

XX The present invention describes a method (M1) for screening for an

CC anti-neoplastic agent. The method involves exposing cells to a chemical

CC agent to be tested for anti-neoplastic activity, determining a change in

CC expression of at least one gene (I) of a signature gene set, where (I)

CC comprises a sequence (S) selected from 8447 sequences (given in ABL61664

CC to ABL70110), or is at least 95% identical to (S), where a change in

CC expression is indicative of anti-neoplastic activity. (I) has cytostatic

CC activity and can be used in gene therapy. M1 can be used for screening

CC an anti-neoplastic agent, and can be used for producing a product which

CC is the data collected with respect to the anti-neoplastic agent as a

CC result of M1, and the data is sufficient to convey the chemical

CC structure and/or properties of the agent. M1 can be used in the

CC treatment of cancer such as colon, breast, stomach, lung, thyroid,

CC esophageal, ovarian, kidney, prostate or pancreatic cancer,

CC adenocarcinoma, carcinoma, clear cell cancer, infiltrating ductal cancer,

CC infiltrating lobular cancer, squamous cell carcinoma, neuroendocrine

CC carcinoma, papillary carcinoma and Wilms' tumour.

XX Sequence 405 BP; 88 A; 97 C; 98 G; 121 T; 1 other;

SQ

Query Match 100.0%; Score 31; DB 24; Length 405;

Best Local Similarity 100.0%; Pred. No. 0.00069;

Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATCATGCTCCCTCTGGCTATCATGCCAACT 31

|||||

Db 384 ATCATGCTCCCTCTGGCTATCATGCCAACT 354

RESULT 2

AAN60429

ID AAN60429 standard; cDNA; 425 BP.

XX AAN60429;

AC AAN60429;

XX 26-JUN-1991 (first entry)

DT Sequence encoding human inhibin B subunit.

XX Hormone; inhibin agonist; antagonist; reproductive; gonad; ss.

XX Homo sapiens.

OS

XX Key Location/Qualifiers

FT CDS 1..42

FT mat_peptide 43..393

FT /*tag= a

FT /*tag= b

XX W08606076-A.

XX 23-OCT-1986.

XX 14-APR-1986; 85WO-AU000097.

XX 20-DEC-1985; 85AU-0003961.

XX 18-APR-1985; 85AU-0000194.

XX 06-SEP-1985; 85AU-0002320.

XX 29-OCT-1985; 85AU-0003157.

XX 19-DEC-1985; 85AU-0003960.

XX 01-JAN-1986; 86AU-0059039.

XX 02-APR-1987; 87AU-0071015.

XX 05-MAY-1986; 86CN-0103459.

XX (BIOT-) BIOTECHN AUSTR PTY.

XX (MONU) MONASH UNIV.

XX (HENR) PRICE HENRY'S HOSPITAL.

XX (SVIN) ST VINCENTS'S INST MED RE.

XX Forage R, Stewart A, Robertson D, Dekretser DM;

XX WPI: 1986-291647/44.

XX P-PSDB; AAP60520.

XX New polynucleotide sequences and recombinant DNA - encoding

PT inhibin and synthetic peptides useful for affecting gonadal

PT function

XX Claim 8; Fig 8; 71pp; English.

XX DNA encoding inhibin and inhibin or part, analogues, homologues or

CC precursors thereof when produced by recombinant techniques are also

CC claimed, as well as pharmaceutical compositions thereof. These may

CC be used as an inhibin agonist, antagonist or for eliciting an

CC antigenic response to affect gonadal function or reproductive

CC physiology.

XX Sequence 425 BP; 103 A; 116 C; 115 G; 91 T; 0 other;

SQ

Query Match 100.0%; Score 31; DB 7; Length 425;

Best Local Similarity 100.0%; Pred. No. 0.00069;

Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATCATGCTCCCTCTGGCTATCATGCCAACT 31

|||||

Db 127 ATCATGCTCCCTCTGGCTATCATGCCAACT 157

RESULT 3

AAH57530
ID AAH57530 standard; cDNA; 1620 BP.
XX
AC AAH57530;
XX
DT 10-SEP-2001 (first entry)
XX
DE Human pancreas cell specific cDNA sequence SEQ ID NO:370.
XX
KW Human; tissue specific; diagnosis; brain; heart; skeletal muscle;
KW lung; liver; uterus; ovary; stomach; intestine; kidney; pancreas; ss;
KW metabolic disease; developmental disease; cytostatic; immunomodulatory;
KW neuroprotective; gene therapy; cancer; immunopathology; neuropathology.
XX
OS Homo sapiens.
XX
PN WC200132927-A2.
XX
PD 10-MAY-2001.
XX
PF 02-NOV-2000; 2000WO-US30396.
XX
PR 04-NOV-1999; 99US-0163508.
XX
PA (INCY-) INCYTE GENOMICS INC.
XX
PI Sornasse T, Seilhamer JJ, Watson GA;
XX
DR WPI: 2001-291057/30.
XX
PT New cell and tissue specific polynucleotides useful for diagnosis.
PT prognosis or monitoring of treatments for disorders where the gene is
PT associated with a cancer, immunopathology or neuropathology -
XX
PS Claim 1; Page 287-288; 327pp; English.
XX
CC AAH57161 to AAH57576 represent cell and tissue specific polynucleotide
CC sequences (I). (I) can have cytostatic, immunomodulatory and
CC neuroprotective activities, and can be used in gene therapy. (I) and
CC proteins (II) encoded by then are used in high throughput screening
CC assays to select DNA molecules, RNA molecules, peptide nucleic acids,
CC mimetics, peptides, proteins, agonists, antagonists, antibodies or
CC their fragments, immunoglobulins, inhibitors, drug compounds and
CC pharmaceutical agents. Expression of (I) in a sample indicates the
CC differentiation of embryonic stem cells into a tissue selected from
CC brain, heart, kidney, liver, lung, skeletal muscle or pancreatic
CC tissues. (I) and (II) are used to produce an expression profile that
CC defines a metabolic or developmental process, treatment, condition,
CC disease or disorder. The gene profile can be used for diagnosis,
CC prognosis or monitoring of treatments and for investigating a
CC predisposition to a disorder where the gene is associated with a
CC cancer, immunopathology or neuropathology.
XX
SQ Sequence 1620 BP; 475 A; 377 C; 476 G; 291 T; 1 other;
Query Match 100.0%; Score 31; DB 22; Length 1620;
Best Local Similarity 100.0%; Pred. No. 0.00086;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATCATGTGCTCCCTGCTATCATGCCAACT 31
Dd 1230 ATCATGTGCTCCCTGCTATCATGCCAACT 1260
RESULT 4
AAH70315
ID AAN70315 standard; cDNA; 1630 BP.
XX
AC AAN70315;
XX
DT 09-APR-1991 (first entry)
XX
DE Sequence encoding human inhibin beta-chain precursor beta-A.

XX
KW Fertility control; contraception; hormone; spermatogenesis; ss.
XX
OS Homo sapiens.
XX
FH Key location/Qualifiers
FT 37..320 /*tag= a
FT CDS /product=signal sequence
FT 321..1166 /*tag= b
FT CDS /product=pro region
FT mat_peptide 1167..11517
FT /*tag= c
XX
PN EP22491-A.
XX
PD 20-MAY-1987.
XX
PF 02-OCT-1986; 86EP-0307586.
XX
PR 12-SEP-1986; 86US-0906729.
PR 03-OCT-1985; 85US-0783910.
PR 10-FEB-1986; 86US-0827710.
XX
PA (GETH) GENENTECH INC.
XX
PI Mason AJ, Seeburg PH;
XX
DR WPI: 1987-137512/20.
DR P-PSDB; AAF70203.
XX
PT Recombinant human or porcine inhibin or activin - used for
PT modulating clinical condition or reproductive physiology of
PT animals.
XX
PS Disclosure; Fig 8A; 48pp; English.
XX
CC A compn. comprising human or porcine inhibin which is completely
CC free of unidentified or porcine proteins is claimed. Also claimed
CC are non chromosomal DNA encoding inhibin-alpha or an inhibin-beta
CC chain. Sequencing of inhibin-encoding cDNA has led to the
CC identification of prodomain regions located N terminal to the
CC mature inhibin chains that represent coordinately expressed
CC biologically active polypeptides. The prodomain regions or
CC prodomain immunogens are useful in monitoring preproinhibin
CC processing in transformant cell culture or in experiments directed
CC at modulating the clinical cond. or reproductive physiology of
CC animals.
XX
SQ Sequence 1630 BP; 472 A; 390 C; 466 G; 302 T; 0 other;
Query Match 100.0%; Score 31; DB 8; Length 1630;
Best Local Similarity 100.0%; Pred. No. 0.00086;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATCATGTGCTCCCTGCTATCATGCCAACT 31
Dd 1251 ATCATGTGCTCCCTGCTATCATGCCAACT 1281
RESULT 5
AAS06008
ID AAS06008 standard; DNA; 1700 BP.
XX
AC AAS06008;
XX
DT 12-SEP-2001 (first entry)
XX
DE Angiotensin converting enzyme (ACEV) splice variant DNA #8.
XX
KW Angiotensin converting enzyme splice variant; ACEV; interleukin 6;
KW granulocyte colony stimulating factor receptor; glucagon; hypertrophy;

KW platelet-derived endothelial cell growth factor; cardiovascular disease;
 KW cellular tumour antigen P53; cyclin-dependent kinase inhibitor 1C; ds;
 KW vasoactive intestinal polypeptide receptor 2; arteriosclerosis; cancer;
 KW myocardial infarction; coronary arterial thrombosis; renal disease;
 KW diabetic nephropathy; muscular disease; immune disorder; sarcoidosis;
 KW multiple sclerosis; immune complex nephritis; deep vein thrombosis;
 KW nonaroidotic pulmonary granulomatous disease; endothelial abnormality;
 KW vascular disorder; asbestosis.
 XX Homo sapiens.
 XX WO200136632-A2.
 XX PD 25-MAY-2001.
 XX
 XX PF 17-NOV-2000; 2000WO-1L00766.
 XX
 XX PR 17-NOV-1999; 99IL-0132978.
 XX PR 10-DEC-1999; 99IL-0133455.
 XX
 XX PA (COMP-) COMPUEN LTD.
 XX
 XX PI Levine Z, David A, Azar I, Khosravi R, Bernstein J;
 XX
 XX DR WPI; 2001-336004/35.
 XX DR P-PSDB; RAU02908.
 XX
 XX PT Novel alternative splicing variants e.g. variant of angiotensin
 XX PT converting enzyme (ACEV), useful in identifying candidate compounds
 XX PT capable of binding to the variant and to detect anti-variant antibodies
 XX PT
 XX PS Claim 1; Page 319; 519pp; English.
 XX
 CC The sequence represents a DNA encoding an angiotensin converting enzyme
 CC splice variant (ACEV) polypeptide. The polypeptides of the invention
 CC include variants of granulocyte colony stimulating factor receptor,
 CC glucagon, interleukin 6, platelet-derived endothelial cell growth factor,
 CC cyclin-dependent kinase inhibitor 1C, cellular tumour antigen P53, and
 CC vasoactive intestinal polypeptide receptor 2. The polypeptides and their
 CC associated nucleic acids are useful for identification of variant
 CC sequences and detection of candidate compounds capable of binding the
 CC molecules. The sequences of the invention can be used in the treatment
 CC and diagnosis of various disorders including cardiovascular diseases such
 CC as arteriosclerosis, myocardial infarction and coronary arterial
 CC thrombosis, renal diseases such as diabetic nephropathy, muscular
 CC diseases such as hypertrophy, immune disorders such as immune complex
 CC nephritis, multiple sclerosis, cancer, sarcoidosis, nonaroidotic
 CC pulmonary granulomatous diseases such as asbestosis and vascular
 CC pathologies involving an endothelial abnormality such as deep vein
 CC thrombosis.
 XX
 XX SQ Sequence 1700 BP; 509 A; 387 C; 440 G; 362 T; 2 other;
 Query Match 100.0%; Score 31; DB 22; Length 1700;
 Best Local Similarity 100.0%; Pred. No. 0.00087;
 Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATCATGCTCCCTCGCTATCATGCCAACT 31
 Db 820 ATCATGCTCCCTCGCTATCATGCCAACT 850
 RESULT 6
 AAQ01648
 ID AAQ01648 standard; DNA; 1873 BP.
 XX
 XX AC AAQ01648;
 XX
 XX DT 27-JUL-1990 (first entry)
 XX
 XX DE BUF-3 gene for human differentiation inducing factor.
 XX

KW BUF-3; dhfr; dihydrofolic acid reductase; differentiation.
 XX Homo sapiens.
 XX
 XX FH Location/Qualifiers
 XX CDS 119..11397
 XX FT /*tag= a
 XX FT 983..1397
 XX FT /*tag= b
 XX FT /label=BUF-3 subunit
 XX
 XX JP02009388-A.
 XX PN
 XX PD 12-JAN-1990.
 XX
 XX PF 08-JUL-1988; 88JP-0170142.
 XX
 XX PR 09-MAR-1988; 88JP-0055270.
 XX
 XX PA (AJIN) AJINOMOTO KK.
 XX
 XX WPI; 1990-055348/08.
 XX DR P-PSDB; AAR05413.
 XX
 XX PT Physiologically active protein prepn. -
 XX PT by transforming plasmid having gene coding physiologically
 XX PT active protein and gene of dihydrofolic acid reductase to hamster
 XX PT ovary etc.
 XX
 XX PS Example 1; Fig 1; 12pp; Japanese.
 XX
 CC Gene may be expressed by transforming a dhfr negative strain of CHO cells
 CC with an active BUF-3 gene and dhfr carrying vector. The BUF-3 gene is
 CC a cell differentiating factor.
 XX
 XX SQ Sequence 1873 BP; 566 A; 431 C; 520 G; 356 T; 0 other;
 Query Match 100.0%; Score 31; DB 11; Length 1873;
 Best Local Similarity 100.0%; Pred. No. 0.00088;
 Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ATCATGCTCCCTCGCTATCATGCCAACT 31
 Db 1133 ATCATGCTCCCTCGCTATCATGCCAACT 1163
 RESULT 7
 ABL62912
 ID ABL62912 standard; DNA; 4068 BP.
 XX
 XX AC ABL62912;
 XX
 XX DT 15-MAY-2002 (first entry)
 XX
 XX DE Breast cancer related gene sequence SEQ ID NO:1249.
 XX
 XX KW Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
 XX KW stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
 XX KW cytostatic; gene therapy; antineoplastic; Wilm's tumour; adenocarcinoma;
 XX KW gene; ds.
 XX
 XX OS Homo sapiens.
 XX
 XX WO200194629-A2.
 XX PN
 XX PD 13-DEC-2001.
 XX
 XX PF 30-MAY-2001; 2001WO-US10838.
 XX
 XX PR 05-JUN-2000; 2000US-209473P.
 XX PR 05-JUN-2000; 2000US-209531P.
 XX PR 18-SEP-2000; 2000US-233133P.
 XX PR 18-SEP-2000; 2000US-233617P.


```

XX PA (AVAL-) AVALON PHARM.
XX
XX PI Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;
PI Soppet DR, Weaver Z;
XX
XX DR WPI; 2002-188264/24.
XX
XX PT Screening for anti-neoplastic agent involves exposing cells to a
PT chemical agent to be tested for anti-neoplastic activity, and
PT determining a change in expression of a gene of a signature gene set -
XX
XX PS Claim 1; SEQ ID 1454; 44pp; English.
XX
XX CC The present invention describes a method (M1) for screening for an
XX anti-neoplastic agent. The method involves exposing cells to a chemical
XX agent to be tested for anti-neoplastic activity, determining a change in
XX expression of at least one gene (I) of a signature gene set, where (I)
XX comprises a sequence (S) selected from 8447 sequences (given in ABL61664
XX to ABL70110), or is at least 95% identical to (S), where a change in
XX expression is indicative of anti-neoplastic activity. (I) has cytostatic
XX activity and can be used in gene therapy. M1 can be used for screening
XX an anti-neoplastic agent, and can be used for producing a product which
XX is the data collected with respect to the anti-neoplastic agent as a
XX result of M1, and the data is sufficient to convey the chemical
XX structure and/or properties of the agent. M1 can be used in the
XX treatment of cancer such as colon, breast, stomach, lung, thyroid,
XX oesophageal, ovarian, kidney, prostate or pancreatic cancer,
XX adenocarcinoma, carcinoma, clear cell cancer, infiltrating ductal cancer,
XX infiltrating lobular cancer, squamous cell carcinoma, neuroendocrine
XX carcinoma, papillary carcinoma and Wilm's tumour.
XX
XX SQ Sequence 4068 BP; 1291 A; 744 C; 893 G; 1140 T; 0 other;
XX
XX Query Match 100.0%; Score 31; DB 24; Length 4068;
XX Best Local Similarity 100.0%; Pred. No. 0.001;
XX Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 ATCATTCCTCCCTCGCTATCATGCCCACT 31
XX |||||||
XX DB 1093 ATCATTCCTCCCTCGCTATCATGCCCACT 1123
XX
XX RESULT 9
XX ABL64417
XX ID ABL64417 standard; DNA; 4068 BP.
XX
XX AC ABL64417;
XX
XX DT 15-MAY-2002 (first entry)
XX
XX DE Stomach cancer related gene sequence SEQ ID NO:2754.
XX
XX KW Human; cancer; colon; breast; ovary; oesophagus; kidney; thyroid;
XX stomach; lung; prostate; pancreas; carcinoma; antitumour; cancerous;
XX cytostatic; gene therapy; antineoplastic; Wilm's tumour; adenocarcinoma;
XX gene; ds.
XX
XX OS Homo sapiens.
XX
XX PN W0200194629-A2.
XX
XX PD 13-DEC-2001.
XX
XX PF 30-MAY-2001; 2001WO-US10838.
XX
XX PR 05-JUN-2000; 2000US-209473P.
XX PR 05-JUN-2000; 2000US-209531P.
XX PR 18-SEP-2000; 2000US-233133P.
XX PR 18-SEP-2000; 2000US-233617P.
XX PR 20-SEP-2000; 2000US-234009P.
XX PR 20-SEP-2000; 2000US-234034P.
XX PR 20-SEP-2000; 2000US-234052P.
XX

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PR 27-SEP-2000; 2000US-235720P.
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PR 28-SEP-2000; 2000US-236032P.
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PR 28-SEP-2000; 2000US-236109P.
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PR 29-SEP-2000; 2000US-236891P.
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PR 02-OCT-2000; 2000US-237173P.
PR 02-OCT-2000; 2000US-237278P.
PR 02-OCT-2000; 2000US-237294P.
PR 02-OCT-2000; 2000US-237295P.
PR 02-OCT-2000; 2000US-237316P.
PR 03-OCT-2000; 2000US-237425P.
PR 03-OCT-2000; 2000US-237598P.
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PR 03-OCT-2000; 2000US-237606P.
PR 03-OCT-2000; 2000US-237608P.
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PR 01-NOV-2000; 2000US-245084P.
XX (AVAL-) AVALON PHARM.
XX
XX PI Young PE, Augustus M, Carter KC, Ebner R, Endress G, Horrigan S;
XX Soppet DR, Weaver Z;
XX
XX DR WPI; 2002-188264/24.
XX
XX PT Screening for anti-neoplastic agent involves exposing cells to a
XX chemical agent to be tested for anti-neoplastic activity, and
XX determining a change in expression of a gene of a signature gene set -
XX
XX PS Claim 1; SEQ ID 2754; 44pp; English.
XX
XX CC The present invention describes a method (M1) for screening for an
XX anti-neoplastic agent. The method involves exposing cells to a chemical
XX agent to be tested for anti-neoplastic activity, determining a change in
XX expression of at least one gene (I) of a signature gene set, where (I)
XX comprises a sequence (S) selected from 8447 sequences (given in ABL61664
XX to ABL70110), or is at least 95% identical to (S), where a change in
XX expression is indicative of anti-neoplastic activity. (I) has cytostatic
XX activity and can be used in gene therapy. M1 can be used for screening
XX an anti-neoplastic agent, and can be used for producing a product which
XX is the data collected with respect to the anti-neoplastic agent as a
XX result of M1, and the data is sufficient to convey the chemical
XX structure and/or properties of the agent. M1 can be used in the
XX treatment of cancer such as colon, breast, stomach, lung, thyroid,
XX oesophageal, ovarian, kidney, prostate or pancreatic cancer,
XX adenocarcinoma, carcinoma, clear cell cancer, infiltrating ductal cancer,
XX infiltrating lobular cancer, squamous cell carcinoma, neuroendocrine
XX carcinoma, papillary carcinoma and Wilm's tumour.
XX
XX SQ Sequence 4068 BP; 1291 A; 744 C; 893 G; 1140 T; 0 other;
XX
XX Query Match 100.0%; Score 31; DB 24; Length 4068;
XX Best Local Similarity 100.0%; Pred. No. 0.001;
XX Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY Match
XX Best Local Similarity 100.0%;
XX Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX

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PR 17-NOV-2000; 2000US-0249299.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250160.
PR 01-DEC-2000; 2000US-0250391.
PR 05-DEC-2000; 2000US-0251030.
PR 05-DEC-2000; 2000US-0251388.
PR 05-DEC-2000; 2000US-0256719.
PR 06-DEC-2000; 2000US-0251479.
PR 08-DEC-2000; 2000US-0251856.
PR 08-DEC-2000; 2000US-0251868.
PR 08-DEC-2000; 2000US-0251869.
PR 08-DEC-2000; 2000US-0251989.
PR 08-DEC-2000; 2000US-0251990.
PR 11-DEC-2000; 2000US-0254097.
PR 05-JAN-2001; 2001US-0259678.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Barash SC, Ruben SM;
XX
XX WPI; 2001-465570/50.
XX
XX Isolated nucleic acid molecule encoding a reproductive system antigen
XX is used in preventing, treating or ameliorating a medical condition -
XX
XX Disclosure; SEQ ID NO 8179; 1297pp + Sequence Listing; English.
XX
XX The present invention provides the protein and coding sequences of a
XX number of human reproductive system related antigens. These can be used
XX in the prevention and treatment of reproductive system disorders,
XX including cancer. The present sequence is a genomic sequence encoding a
XX protein of the invention.
XX
XX Sequence 14416 BP; 4206 A; 3105 C; 3196 G; 3909 T; 0 other;
XX
XX Query Match 100.0%; Score 31; DB 22; Length 14416;
XX Best Local Similarity 100.0%; Pred. No. 0.0012;
XX Matches 31; Conservative 0; Mismatches 0; Gaps 0;
XX
XX 1 ATCATTGCTCCCTGGCTATCATGCCAACT 31
XX |||||||
Db 13179 ATCATTGCTCCCTGGCTATCATGCCAACT 13209
XX
XX
XX RESULT 11
XX ABL98344
XX ID ABL98344 standard; DNA; 14416 BP.
XX
XX ABL98344;
XX
XX 21-JUN-2002 (first entry)
XX
XX Human testicular antigen encoding DNA fragment SEQ ID NO: 2996.
XX
XX Human; testicular antigen; testes; cancer; metastasis; immune disorder;
XX reproductive system disorder; urinary system disorder; gene therapy;
XX cardiovascular disorder; respiratory disorder; neurological disorder;
XX gastrointestinal disease; infection; cytostatic; gene; ds.
XX
XX Homo sapiens.
XX
XX W0200155317-A2.
XX
XX 02-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US01329.
XX
XX 31-JAN-2000; 2000US-0179065.
XX 04-FEB-2000; 2000US-0180628.
XX 24-FEB-2000; 2000US-0184664.
XX 02-MAR-2000; 2000US-0186350.
XX 16-MAR-2000; 2000US-0189474.
PR 17-MAR-2000; 2000US-0190076.
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PR 26-JUL-2000; 2000US-0220964.
PR 14-AUG-2000; 2000US-0224518.
PR 14-AUG-2000; 2000US-0224519.
PR 14-AUG-2000; 2000US-0225213.
PR 14-AUG-2000; 2000US-0225214.
PR 14-AUG-2000; 2000US-0225266.
PR 14-AUG-2000; 2000US-0225267.
PR 14-AUG-2000; 2000US-0225268.
PR 14-AUG-2000; 2000US-0225270.
PR 14-AUG-2000; 2000US-0225447.
PR 14-AUG-2000; 2000US-0225757.
PR 14-AUG-2000; 2000US-0225758.
PR 14-AUG-2000; 2000US-0225759.
PR 18-AUG-2000; 2000US-0226279.
PR 22-AUG-2000; 2000US-0226681.
PR 22-AUG-2000; 2000US-0226868.
PR 22-AUG-2000; 2000US-0227182.
PR 23-AUG-2000; 2000US-0227009.
PR 30-AUG-2000; 2000US-0228924.
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PR 01-SEP-2000; 2000US-0229343.
PR 01-SEP-2000; 2000US-0229344.
PR 01-SEP-2000; 2000US-0229345.
PR 05-SEP-2000; 2000US-0229509.
PR 05-SEP-2000; 2000US-0229513.
PR 06-SEP-2000; 2000US-0230437.
PR 06-SEP-2000; 2000US-0230438.
PR 08-SEP-2000; 2000US-0231242.
PR 08-SEP-2000; 2000US-0231243.
PR 08-SEP-2000; 2000US-0231244.
PR 08-SEP-2000; 2000US-0231413.
PR 08-SEP-2000; 2000US-0231414.
PR 08-SEP-2000; 2000US-0232080.
PR 08-SEP-2000; 2000US-0232081.
PR 12-SEP-2000; 2000US-0231968.
PR 14-SEP-2000; 2000US-0232397.
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PR 25-SEP-2000; 2000US-0234997.
PR 25-SEP-2000; 2000US-0234998.
PR 26-SEP-2000; 2000US-0235484.
PR 27-SEP-2000; 2000US-0235834.
PR 27-SEP-2000; 2000US-0235836.
PR 29-SEP-2000; 2000US-0236327.
PR 29-SEP-2000; 2000US-0236367.
PR 29-SEP-2000; 2000US-0236368.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236370.
PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
PR 02-OCT-2000; 2000US-0237038.
PR 02-OCT-2000; 2000US-0237039.
PR 02-OCT-2000; 2000US-0237040.
PR 13-OCT-2000; 2000US-0239935.
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PR 13-OCT-2000; 2000US-0239937.
 PR 20-OCT-2000; 2000US-0240960.
 PR 20-OCT-2000; 2000US-0241221.
 PR 20-OCT-2000; 2000US-0241785.
 PR 20-OCT-2000; 2000US-0241786.
 PR 20-OCT-2000; 2000US-0241787.
 PR 20-OCT-2000; 2000US-0241808.
 PR 20-OCT-2000; 2000US-0241809.
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 PR 01-NOV-2000; 2000US-0244617.
 PR 08-NOV-2000; 2000US-0246474.
 PR 08-NOV-2000; 2000US-0246475.
 PR 08-NOV-2000; 2000US-0246476.
 PR 08-NOV-2000; 2000US-0246477.
 PR 08-NOV-2000; 2000US-0246478.
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 PR 08-NOV-2000; 2000US-0246611.
 PR 08-NOV-2000; 2000US-0246613.
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 PR 17-NOV-2000; 2000US-0249208.
 PR 17-NOV-2000; 2000US-0249209.
 PR 17-NOV-2000; 2000US-0249210.
 PR 17-NOV-2000; 2000US-0249211.
 PR 17-NOV-2000; 2000US-0249212.
 PR 17-NOV-2000; 2000US-0249213.
 PR 17-NOV-2000; 2000US-0249214.
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 PR 17-NOV-2000; 2000US-0249216.
 PR 17-NOV-2000; 2000US-0249217.
 PR 17-NOV-2000; 2000US-0249218.
 PR 17-NOV-2000; 2000US-0249244.
 PR 17-NOV-2000; 2000US-0249245.
 PR 17-NOV-2000; 2000US-0249264.
 PR 17-NOV-2000; 2000US-0249265.
 PR 17-NOV-2000; 2000US-0249267.
 PR 17-NOV-2000; 2000US-0249297.
 PR 17-NOV-2000; 2000US-0249299.
 PR 17-NOV-2000; 2000US-0249300.
 PR 01-DEC-2000; 2000US-0250160.
 PR 01-DEC-2000; 2000US-0250391.
 PR 05-DEC-2000; 2000US-0251030.
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 PR 05-DEC-2000; 2000US-0256719.
 PR 06-DEC-2000; 2000US-0251479.
 PR 08-DEC-2000; 2000US-0251856.
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 PR 08-DEC-2000; 2000US-0251869.
 PR 08-DEC-2000; 2000US-0251869.
 PR 08-DEC-2000; 2000US-0251989.
 PR 11-DEC-2000; 2000US-0251990.
 PR 11-DEC-2000; 2000US-0254097.
 PR 05-JAN-2001; 2001US-0259678.
 PA (HUMA-) HUMAN GENOME SCI INC.
 XX
 PT Rosen CA, Barash SC, Ruben SM;
 DR WPI; 2001-483232/52.
 XX
 PT Nucleic acids encoding 973 human testicular antigen polypeptides,
 PT useful for preventing, diagnosing and/or treating testicular cancer
 XX
 PS Disclosure; SEQ ID NO 2996; 766pp; English.
 XX
 XX The present invention provides the protein and coding sequences of 973
 CC human testicular antigens, and fragments of their genomic sequences. The
 CC sequences can be used in the treatment of cardiovascular, urinary system,

CC reproductive system, immune, respiratory, neurological and
 CC gastrointestinal disorders, infections, and particularly cancer,
 CC especially testicular cancers. The present sequence is a DNA encoding a
 CC protein fragment of the invention.
 XX
 SQ Sequence 14416 BP; 4206 A; 3105 C; 3196 G; 3909 T; 0 other;
 Query Match 100.0%; Score 31; DB 23; Length 14416;
 Best local similarity 100.0%; Pred. No. 0.0012;
 Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 ATCATTGCTCCCTGCTATCATGCCAACT 31
 Db 13179 ATCATTGCTCCCTGCTATCATGCCAACT 13209
 RESULT 12
 ID AAN60427 standard; cDNA; 958 BP.
 AC AAN60427;
 XX
 XX 26-JUN-1991 (first entry)
 DE Sequence encoding bovine inhibin B subunit.
 KW Hormone; inhibin agonist; antagonist; reproductive; gonad; ss.
 OS Bos taurus.
 XX
 FH Key Location/Qualifiers
 FT CDS 1..516
 FT mat_peptide /*tag= a
 FT /*tag= b
 PN WO8606076-A.
 XX
 PD 23-OCT-1986.
 XX
 PF 14-APR-1986; 86WO-AU000097.
 XX
 PR 20-DEC-1985; 85AU-0003961.
 PR 18-APR-1985; 85AU-0000194.
 PR 06-SEP-1985; 85AU-0002320.
 PR 29-OCT-1985; 85AU-0003157.
 PR 19-DEC-1985; 85AU-0003960.
 PR 01-JAN-1986; 86AU-0059039.
 PR 02-APR-1987; 87AU-0071015.
 PR 05-MAY-1986; 86CN-0103459.
 XX
 PA (BIOT-) BIOTECHN AUSTR PTY.
 PA (MONU) MONASH UNIV.
 PA (HENR-) PRICE HENRY'S HOSPITAL.
 PA (SVIN-) ST VINCENT'S S INST MED RE.
 XX
 PI Forage R, Stewart A, Robertson D, Dekretser DM;
 XX
 DR WPI; 1986-291647/44.
 DR P-PSDB; AAP60518.
 XX
 PT New polynucleotide sequences and recombinant DNA - encoding
 PT inhibin and synthetic peptides useful for affecting gonadal
 PT function
 XX
 PS Claim 8; Fig 6; 71pp; English.
 XX
 CC DNA encoding inhibin and inhibin or part, analogues, homologues or
 CC precursors thereof when produced by recombinant techniques are also
 CC claimed, as well as pharmaceutical compositions thereof. These may
 CC be used as an inhibin agonist, antagonist or eliciting an
 CC antigenic response to affect gonadal function or reproductive
 CC physiology.

XX SQ Sequence 958 BP; 242 A; 240 C; 302 G; 174 T; 0 other;
 Query Match 84.5%; Score 26.2; DB 7; Length 958;
 Best Local Similarity 90.3%; Pred. No. 0.094;
 Matches 28; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 ATCATGTCCTCTCGGCTATCATGCCAACT 31
 ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 601 ATCATGTCCTCTCGGCTATCATGCCAACT 631

RESULT 13
 AAN70317
 ID AAN70317 standard; cDNA; 3588 BP.
 XX
 AC
 XX
 DT 09-APR-1991 (first entry)
 XX
 DE Sequence encoding porcine inhibin beta-chain precursor beta-A.
 XX
 KW Fertility control; contraception; hormone; spermatogenesis; ss.
 XX
 OS Sus scrofa domestica.
 XX
 FH Key Location/Qualifiers
 CDS 34..957 /tag= a
 FT /product= hydrophobic signal sequence a pro-region
 FT mat_peptide 958..1307
 FT /tag= b
 FT polyA_signal 3551..3556
 FT /tag= c
 XX
 PN EP222491-A.
 XX
 XX 20-MAY-1987.
 XX
 XX 02-OCT-1986; 86EP-0307586.
 XX
 PR 12-SEP-1986; 86US-0906729.
 PR 03-OCT-1985; 85US-0783910.
 PR 10-FEB-1986; 86US-0827710.
 XX
 PA (GETH) GENENTECH INC.
 XX
 PI Mason AJ, Seeburg PH;
 XX
 DR WPI: 1987-137512/20.
 DR P-PSDB; AAP70200.
 XX
 XX Recombinant human or porcine inhibin or activin - used for
 PT modulating clinical condition or reproductive physiology of
 PT animals.
 XX
 PS Disclosure; Fig 1B; 48pp; English.
 XX
 XX A compsn. comprising human or porcine inhibin which is completely
 CC free of unidentified or porcine proteins is claimed. Also claimed
 CC are non chromosomal DNA encoding inhibin-alpha or an inhibin-beta
 CC chain. Sequencing of inhibin-encoding cDNA has led to the
 CC identification of prodomain regions located N-terminal to the
 CC mature inhibin chains that represent coordinately expressed
 CC biologically active polypeptides. The prodomain regions or
 CC prodomain immunogens are useful in monitoring preproinhibin
 CC processing in transformant cell culture or in experiments directed
 CC at modulating the clinical condn. or reproductive physiology of
 CC animals.
 XX
 SQ Sequence 3588 BP; 881 A; 986 C; 998 G; 723 T; 0 other;

Query Match 74.2%; Score 23; DB 8; Length 3588;

Best Local Similarity 83.9%; Pred. No. 2.8;
 Matches 26; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 ATCATGTCCTCTCGGCTATCATGCCAACT 31
 ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 1042 ATCATGTCCTCTCGGCTATCATGCCAACT 1072

RESULT 14
 AAQ10890
 ID AAQ10890 standard; DNA; 1667 BP.
 XX
 AC
 XX
 DT 13-MAY-1991 (first entry)
 XX
 DE Encodes Xenopus Bone Morphogenetic Factor B9.
 XX
 KW BMF; osteoporosis; fracture; cartilage; ss.
 XX
 OS Xenopus laevis.
 XX
 FH Key Location/Qualifiers
 FT mat_peptide 651..1040
 FT /tag= a
 FT /product= BMP B9
 XX
 PN EP416578-A.
 XX
 XX 13-MAR-1991.
 XX
 XX 05-SEP-1990; 90EP-0117079.
 XX
 XX 20-JUL-1990; 90JP-0190774.
 PR 06-SEP-1989; 89JP-0229250.
 XX
 PA (TAKE) TAKEDA CHEMICAL IND KK.
 PA (SCIT-) SCITECH RESEARCH CO.
 XX
 PI Murakami K, Ueno N, Kato Y;
 XX
 DR WPI: 1991-075112/11.
 DR P-PSDB; AAR10990.
 XX
 XX Xenopus laevis bone morphogenetic protein and DNA encoding it -
 PT used in therapy of fracture or osteoporosis
 XX
 PS Claim 5; Fig 2; 28pp; English.
 XX
 CC A Xenopus laevis liver-derived DNA library in Charon 28 vector, was
 CC screened with a rat activin beta-A cDNA probe. Five clones were
 CC isolated, including clone B9. They were subcloned in pUC19 and used
 CC to transform competent E.coli HB101 cells. Transformant E.coli HB101/
 CC pXa-9 coding for the B9 BMP was sequenced. See also AAQ10891-7.
 XX
 SQ Sequence 1667 BP; 557 A; 295 C; 355 G; 450 T; 10 other;

Query Match 59.0%; Score 21.4; DB 12; Length 1667;
 Best Local Similarity 80.6%; Pred. No. 12;
 Matches 25; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 1 ATCATGTCCTCTCGGCTATCATGCCAACT 31
 ||||| ||||| ||||| ||||| ||||| ||||| |||||
 Db 777 ATCATGACCTCTCGGCTATCATGCCAACT 807

RESULT 15
 AAL07086
 ID AAL07086 standard; DNA; 9662 BP.
 XX
 AC AAL07086;
 XX
 DT 21-NOV-2001 (first entry)

XX	Human reproductive system related antigen DNA SEQ ID NO: 9774.	PR	14-SEP-2000;	2000US-0233065.
DE		PR	21-SEP-2000;	2000US-0234223.
XX		PR	21-SEP-2000;	2000US-0234274.
XX		PR	25-SEP-2000;	2000US-0234997.
KW	Human; reproductive system related antigen; reproductive system disorder;	PR	25-SEP-2000;	2000US-0234998.
KW	cancer; gene therapy; ds.	PR	26-SEP-2000;	2000US-0235484.
XX		PR	27-SEP-2000;	2000US-0235834.
OS	Homo sapiens.	PR	27-SEP-2000;	2000US-0235836.
XX		PR	29-SEP-2000;	2000US-0236327.
PN	WO200155320-A2.	PR	29-SEP-2000;	2000US-0236367.
XX		PR	29-SEP-2000;	2000US-0236368.
PD		PR	29-SEP-2000;	2000US-0236370.
XX	02-AUG-2001.	PR	02-OCT-2000;	2000US-0236802.
XX		PR	02-OCT-2000;	2000US-0237037.
XX	17-JAN-2001; 2001WO-US01339.	PR	02-OCT-2000;	2000US-0237038.
PR		PR	02-OCT-2000;	2000US-0237039.
PR	31-JAN-2000; 2000US-0179065.	PR	02-OCT-2000;	2000US-0237040.
PR	04-FEB-2000; 2000US-0180628.	PR	02-OCT-2000;	2000US-0237040.
PR	24-FEB-2000; 2000US-0184664.	PR	02-OCT-2000;	2000US-023935.
PR	02-MAR-2000; 2000US-0186350.	PR	13-OCT-2000;	2000US-023935.
PR	16-MAR-2000; 2000US-0189874.	PR	13-OCT-2000;	2000US-023937.
PR	17-MAR-2000; 2000US-0190076.	PR	20-OCT-2000;	2000US-023937.
PR	18-APR-2000; 2000US-0198123.	PR	20-OCT-2000;	2000US-0240960.
PR	19-MAY-2000; 2000US-0205515.	PR	20-OCT-2000;	2000US-0241221.
PR	07-JUN-2000; 2000US-0209467.	PR	20-OCT-2000;	2000US-0241221.
PR	28-JUN-2000; 2000US-0214886.	PR	20-OCT-2000;	2000US-0241785.
PR	30-JUN-2000; 2000US-0215135.	PR	20-OCT-2000;	2000US-0241786.
PR	07-JUL-2000; 2000US-0216647.	PR	20-OCT-2000;	2000US-0241787.
PR	07-JUL-2000; 2000US-0216880.	PR	20-OCT-2000;	2000US-0241808.
PR	11-JUL-2000; 2000US-0217487.	PR	20-OCT-2000;	2000US-0241809.
PR	11-JUL-2000; 2000US-0217496.	PR	01-NOV-2000;	2000US-0241826.
PR	14-JUL-2000; 2000US-0218290.	PR	08-NOV-2000;	2000US-0246474.
PR	26-JUL-2000; 2000US-0220963.	PR	08-NOV-2000;	2000US-0246475.
PR	26-JUL-2000; 2000US-0220964.	PR	08-NOV-2000;	2000US-0246475.
PR	14-AUG-2000; 2000US-0224518.	PR	08-NOV-2000;	2000US-0246476.
PR	14-AUG-2000; 2000US-0224519.	PR	08-NOV-2000;	2000US-0246477.
PR	14-AUG-2000; 2000US-0225213.	PR	08-NOV-2000;	2000US-0246478.
PR	14-AUG-2000; 2000US-0225214.	PR	08-NOV-2000;	2000US-0246523.
PR	14-AUG-2000; 2000US-0225266.	PR	08-NOV-2000;	2000US-0246524.
PR	14-AUG-2000; 2000US-0225267.	PR	08-NOV-2000;	2000US-0246525.
PR	14-AUG-2000; 2000US-0225268.	PR	08-NOV-2000;	2000US-0246526.
PR	14-AUG-2000; 2000US-0225270.	PR	08-NOV-2000;	2000US-0246527.
PR	14-AUG-2000; 2000US-0225447.	PR	08-NOV-2000;	2000US-0246528.
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